



Growth and physiology of the citrange ‘Fepagro C37 Reck’ rootstock inoculated with arbuscular mycorrhizal fungi¹

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ABSTRACT

The use of arbuscular mycorrhizal fungi (AMF) in the production of rootstocks is a strategy to promote faster plant growth. However, this response depends on the symbiont species. The objective of this study was to evaluate the influence of different species of arbuscular mycorrhizal fungus (AMF) on the development and physiology of citrange ‘Fepagro C37 Reck’ rootstock. Four species of AMF were used in the experiment: *Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum*, and *Acaulospora* sp., and a control treatment. The experiment was arranged in a randomized blocks design, with 5 treatments, 3 replicates, and 10 plants per plot. Every 15 days, height (cm) and stem diameter (mm) of plants were measured in the nursery. At 330 days, the number of leaves, leaf area (cm²), fresh and dry mass of roots and shoot (g/plant), root and shoot reserve content, and AMF colonization were evaluated. The AMF inoculation promoted an accelerated vegetative growth, resulting in greater height and diameter. The fungi also promoted greater absorption of most of the nutrients and increase in all other parameters evaluated. However, no effect was found on the reserve content of the rootstock. Roots were colonized with moderate presence of arbuscules and low presence of vesicles. AMF, regardless of species, influence the development and physiology of the citrange ‘Fepagro C37 Reck’ rootstock.

Keywords: endomycorrhizae; citrus; clonal seedling production; vegetative growth; nutrients; colonization.

RESUMO

Desenvolvimento e fisiologia do porta-enxerto citrangeiro ‘Fepagro C37 Reck’ inoculado com fungos micorrízicos arbusculares

A utilização de fungos micorrízicos arbusculares (FMA) na produção de porta-enxertos é uma alternativa para promover um desenvolvimento mais acelerado às plantas. Entretanto, sua resposta depende das espécies dos simbiontes. O objetivo deste estudo foi avaliar a influência de diferentes espécies de FMA no desenvolvimento e fisiologia do porta-enxerto citrangeiro ‘Fepagro C37 Reck’. Foram utilizadas quatro espécies de FMA: *Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp., bem como um tratamento controle. O delineamento experimental foi de blocos ao acaso de 5 tratamentos, com 3 repetições e 10 plantas por parcela. A cada quinze dias, foi medida a altura (cm) e o diâmetro do caule (mm) das plantas no viveiro. Aos 330 dias, foram medidos o número de folhas, a área foliar (cm²), a massa fresca e seca das raízes e da parte aérea (g/planta), o teor de reservas da raiz e da parte aérea, o conteúdo nutricional da parte aérea e a colonização de FMA. A inoculação de FMA promoveu um crescimento vegetativo acelerado, resultando em maior altura e diâmetro. Além disso, os fungos promoveram às plantas maior absorção da maioria dos nutrientes e em todos os outros parâmetros testados. Contudo, não se observou efeito no teor de reservas do porta-enxerto. As raízes foram colonizadas, com moderada presença de arbúsculos e baixa de vesículas. Os FMA, independentemente da espécie, influenciam no desenvolvimento e na fisiologia do porta-enxerto citrangeiro ‘Fepagro C37 Reck’.

Palavras-chave: endomicorrizas; citros; produção de mudas; desenvolvimento vegetativo; nutrientes; colonização.

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INTRODUCTION

Citrus production in Brazil is characterized by a low genetic diversity of rootstocks, making it vulnerable to new diseases (Fochesato *et al.*, 2007). One way to reduce risks is the use of different species/varieties of rootstocks, with different characteristics (Souza *et al.*, 2005; Schäfer *et al.*, 2006).

To achieve diversification, studies are needed to test different species with rootstock potential and that have, for example, resistance to diseases, tolerance to abiotic stresses, and affinity with scion varieties of market interest. In addition, the performance should be analyzed throughout the production cycle, from clonal seedling formation in the nursery, during the initial growth in the orchard, and during the full fruit production (Zaccheo *et al.*, 2012; Gonzatto *et al.*, 2011).

In the state of Rio Grande do Sul, the rootstocks used show low growth rate at the nursery stage because of the low temperatures in the winter. Therefore, the time required for clonal seedling production in the southern region of the country is longer, increasing the production cost and, thus, losing to other regions of Brazil in the competitive market (Souza *et al.*, 2005).

A rootstock with great potential, which has already been used successfully in some orchards of Rio Grande do Sul, is 'Fepagro C37 Reck' [*Poncirus trifoliata* (L.) Raf. x *Citrus sinensis* (L.) Osbeck.]. This is a hybrid with trifoliolate orange (*P. trifoliata* (L.) Raf.), the most commonly used rootstock in the southern region, and has its desirable characteristics, as well as showing a faster growth in the nursery (Oliveira *et al.*, 2010; Koller, 2013).

An alternative technique for accelerating plant growth is the inoculation with arbuscular mycorrhizal fungi (AMF). These are organisms that are symbiotically associated with plant roots (Jayme & Quigley, 2014; Wu *et al.*, 2011a). In this symbiosis, the plant supplies the fungus with energy for growth and reproduction via photosynthesis, and the fungus provides the plant with greater absorption of nutrients beyond the root zone, especially phosphorus (Smith & Smith, 2011). Other important effects of AMF are the increased plant resistance to attack of pathogens of the root system and water absorption capacity (Goltapeh *et al.*, 2008). Therefore, the symbiosis allows the plant a greater absorption of nutrients and water, which accelerates its growth (Jayme & Quigley, 2014).

According to Ortas (2012), citrus plants are usually highly responsive to AMF because the root system lacks absorbent hairs. However, Smith & Smith (2011) pointed out that the response of the plant to mycorrhization depends on several factors; one of them is related to the species of the arbuscular mycorrhizal fungus inoculated, mainly due to compatibility with the host and/or adaptation to the environment.

Thus, the objective of this study was to evaluate the influence of different species of arbuscular mycorrhizal fungi on the growth and physiology of citrange 'Fepagro C37 Reck' rootstock.

MATERIAL AND METHODS

The study was carried out in two phases. The sowing phase (from sowing and AMF inoculation of plants with 10 cm in height) was carried out in a greenhouse of the Horticulture and Forestry Department (HFD), Campus of the Faculdade de Agronomia of the Universidade Federal do Rio Grande do Sul (UFRGS), Porto Alegre, RS - Brazil (30°29'S and 51°06'W). The nursery phase (from pricking out seedlings to 5-L polyethylene bags to the end of the experiment) was conducted in a greenhouse at the Agronomic Experimental Station of the Faculdade de Agronomia of the Universidade Federal do Rio Grande do Sul (EEA/UFRGS, 30°05'22"S, 51°39'08"W), municipality of Eldorado do Sul, RS - Brazil (Km 146, BR 290).

The seeds of the citrange 'Fepagro C37 Reck' rootstock [*P. trifoliata* (L.) Raf. x *C. sinensis* (L.) Osbeck.] were obtained from mature fruits harvested from mother plants grown in the citrus collection of the Experimental Agronomic Station at UFRGS, using the methodology described by Souza & Schäfer (2006). The seeds were heat treated (soaking in water at 52 °C for 10 minutes).

The AMF inoculants (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum*, and *Acaulospora* sp.) used in the experiment were grown in the HFD Laboratory from an initial inoculum obtained from the Inoculant Bank of the same department. The selected AMF species were selected from an initial study by Focchi *et al.* (2004) who carried out a study on arbuscular mycorrhizal fungi naturally occurring in citrus orchards in Rio Grande do Sul.

The inoculum was produced as follows: four 5L pots were filled with sterilized sand and the inoculum of an AMF species was mixed in the upper 4cm layer of the sand. After inoculation, 20 corn seeds (*Zea mays* L.) were planted per pot. The pots remained in the laboratory under artificial light and hand watered for two months. After this time, the sand was homogenized with the remains of corn roots, and the new inoculum originated from this mixture. Finally, the inoculum was stored in refrigerator for better conservation.

In February 2013, the rootstocks were inoculated with the AMF. The seeds of the rootstock were planted in commercial substrate, consisting of peat, charred rice husk, and vermiculite in plastic trays (60 cm x 40 cm x 10 cm). The substrate was previously autoclaved at 120 °C (1 atm pressure) for one hour, for three times at 24 hour intervals.

The trays were kept on a concrete bench in a protected environment and covered by a protective screen to avoid

insects that could contaminate the experiment. Each tray contained a treatment with AMF (control and the four AMF species).

The trays treated with AMF inoculum were filled with substrate up to the half of the height; and a layer of 400 g of inoculum (approx. 2 mm) was placed on the substrate layer, completing the tray height with substrate. The tray containing the control plants received no inoculum. Finally, 50 seeds of the rootstock were sown per treatment. Trays were hand watered using a hand irrigator, with 1L water per tray every two days.

Seedlings, with an average height of 10 cm (June 8, 2013), were transferred to the greenhouse at the Experimental Agronomic Station of UFRGS in the municipality of Eldorado do Sul, RS. They were pricked out to 5-L polyethylene bags filled with commercial substrate consisting of eucalyptus bark previously disinfested with 7% formaldehyde solution. The seedlings were kept in 80 cm-high wooden benches and received drip irrigation twice a day, 5 minutes each, totaling 50 mL of water volume per day.

The experiment was arranged in a randomized blocks design, with 5 treatments (4 AMF + control), 3 replicates, and 10 plants per plot. Vegetative growth, reserve content, and nutrient content were analyzed in 'Fepagro C37 Reck [*P. trifoliata* (L.) Raf. x *C. sinensis* (L.) Osbeck.] rootstock inoculated with four AMF species (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* and *Acaulospora* sp.) and a control treatment (non-inoculated).

Plant vegetative growth (10 plants per plot) was monitored biweekly, measuring height (cm), from the collar to the apex of the plant with a measuring tape, and stem diameter (mm) with a digital caliper, 1 cm above the substrate level.

The final height (cm), stem diameter (mm), and leaf number per plant of 10 rootstocks per plot were measured 240 days after pricking out the seedlings. Leaf area (cm²) per plant was measured using a leaf area LI-Meter. The fresh root and shoot mass (g) and dry matter of root and shoot (g) were determined after drying the samples in a oven at 65 °C to constant mass.

Dried shoots and roots were ground, mixing 10 plants per treatment and repetition, from which a 1 g sample was collected to evaluate reserve content, according to the method described by Priestley (1965) with modifications. Each sample was wrapped in a filter cloth, forming small bags. The bags were weighed before and after digestion for 8 hours, soaked in aqueous solution containing 5% trichloroacetic acid and 35% methanol, to extract all components of the plant tissue (carbohydrates, fats, etc.) that were not fibers (cellulose, hemicellulose, and lignin). The difference in mass, before and after the digestion,

indicates the percentage of reserves contained. We used 3 replicates for each treatment.

The remaining dry matter from samples of shoots of each treatment, initially ground, was used to determine macronutrients (N, P, K, Ca, Mg, S) and micronutrients (Cu, Zn, Fe, Mn and B) carried out at the Laboratory of Soil and Tissue Analysis of the Faculdade de Agronomia - UFRGS. The results were analyzed according to the Manual of Fertilization and Liming for the States of Rio Grande do Sul and Santa Catarina (SBCS, 2004).

In the same period, two segments of secondary roots were collected from each plant to evaluate the AMF colonization. The roots were washed with distilled water, cut 1 cm in length, and stored in FAA (formaldehyde, alcohol, and acetone) for fixation. Then, root segments were collected randomly, 15 per replicate, totaling 45 segments per treatment, and stained with trypan blue to observe AMF colonization and structures in the roots using a microscope.

After staining, the root segments were mounted on glass slides and examined under a microscope to evaluate the presence and intensity of hyphae, arbuscules, and vesicles. The percentage of colonized roots was calculated based on the number of infected segments in relation to the total material analyzed. Hyphal density was graded by assigning 0 for the absence of structures; 1, for low presence; 2, for moderate presence; and 3, for high presence. The density of vesicles and arbuscules was also graded by a scale from 0 to 3, where the absence of structures was considered as 0; 1 to 50 structures as 1; 51 to 100 as 2; and more than 100 as 3, according to methodology described by Nemec (1992). The process was repeated three times and trained evaluators were used for greater reliability of the results.

Data were analyzed by ANOVA and means were compared by the Duncan test, at the of 1% and 5% of significance level.

RESULTS AND DISCUSSION

The evaluations over the 240 days post-transplant (DAT) to 5-L pots showed that, regardless of treatment, plants had no statistical difference in height up to approximately 130 DAT, (Figure 1).

From that time, the plants inoculated with *Gigaspora margarita*, *Scutellospora heterogama*, and *Glomus etunicatum* showed an exponential primary growth, nearly quintupling the plant height compared to the non-inoculated plants at 240 DAT. Plants inoculated with *Acaulospora* sp. also showed a similar trend, but not as pronounced as with the other species. Non-inoculated plants had a small increase in height after 150 DAT, stabilized in the last days, and reached average of 21 cm at 240 DAT.

The plants of all treatments showed no statistical difference for stem diameter up to 150 days (Figure 2). From this time, the inoculated plants started an exponential secondary growth, similar among all inoculated treatments, reaching twice the diameter compared with the controls at 240 DAT (Figure 2).

There was absence of primary (up to 130 DAT) and secondary (up to 150 DAT) growth, because these periods coincided with the winter in the southern country, with average temperatures ranging from 11.5 °C to 15 °C,

from June to September 2013. For the inoculated plants, besides the effect of temperature, absence of growth in this initial period may have been a consequence of the time required for AMF to colonize the roots and the substrate. Smith & Smith (2011) discussed that AMF need time for hyphae to occupy the volume of the pot and establish their structures in the root tissues of the host. It is only from this period onwards that they provide a significant acceleration of the plant growth, as they will aid nutrient and water absorption.

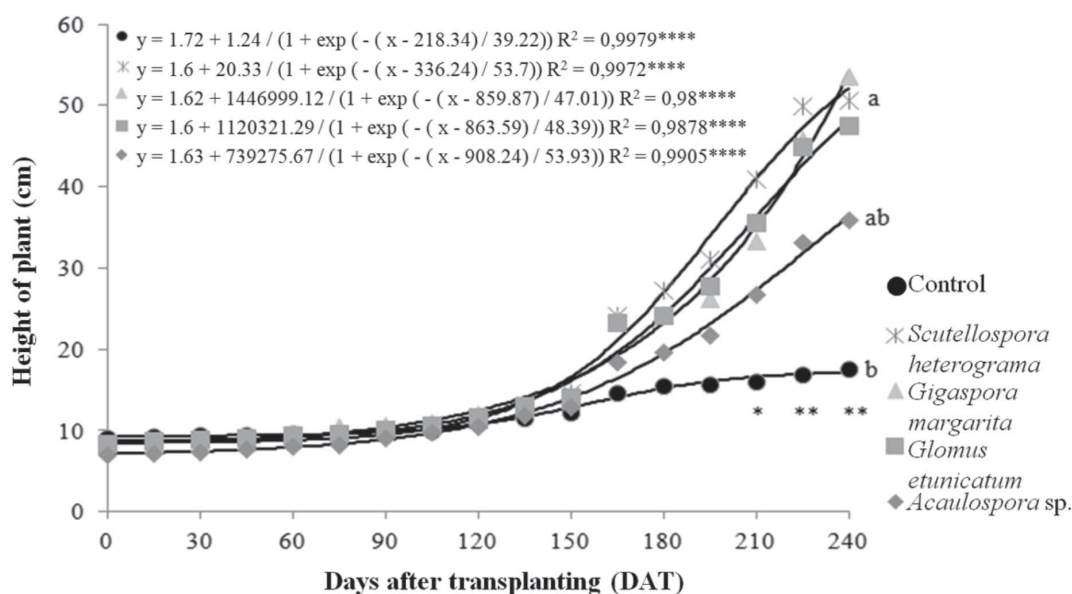


Figure 1: Evolution of the height (cm) of the citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of arbuscular mycorrhizal fungi during 240 days after transplanting in a greenhouse. **** means 0.01% significance by the test F. * and **, mean, respectively, 5% and 1% of significance by the Duncan test. Equal lowercase letters do not differ statistically.

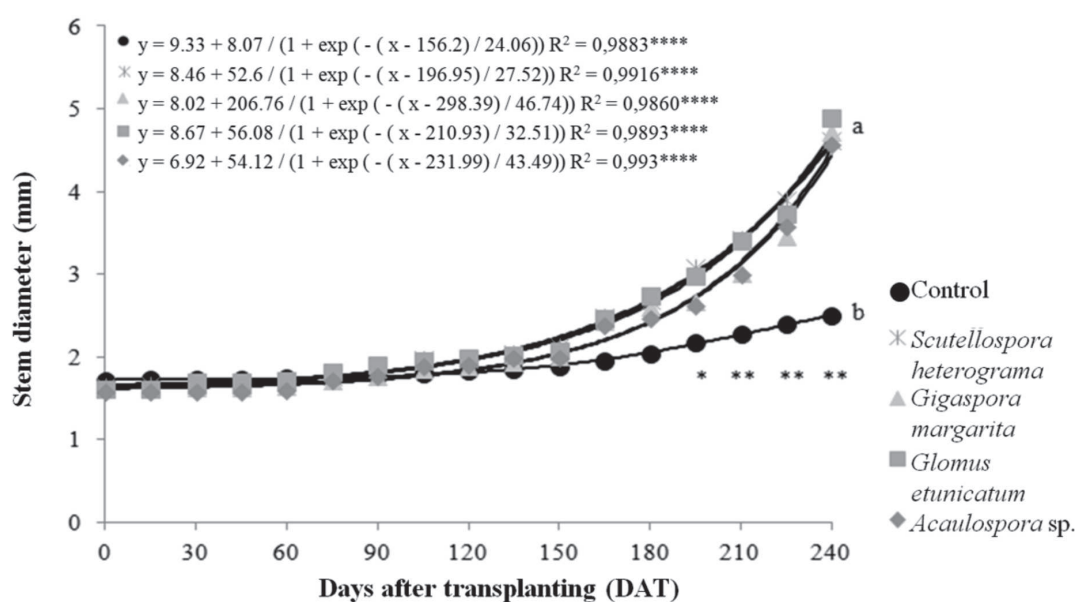


Figure 2: Evolution of stem diameter (mm) of citrange 'Fepagro C37 Reck' rootstock submitted to inoculation of arbuscular mycorrhizal fungi throughout 240 days after transplanting in a greenhouse. **** means 0.01% significance by the test F. * and **, mean, respectively, 5% and 1% of significance by the Duncan test. Equal lowercase letters do not differ statistically.

The acceleration of plant growth provided by AMF results in a reduction in the time to reach the grafting stage, which may reduce production costs of nurseries. It also allows the production of seedlings within 18 to 22 months, required by Normative Instruction n. 24 of December 16, 2005 (Brazil, 2018), sometimes impossible in southern Brazil, due to the low temperatures in autumn/winter.

In addition to increasing height and stem diameter of citrange 'Fepagro C37 Reck' rootstock, inoculation with AMF, independently of the species, doubled the number of leaves, almost tripled the leaf area, and quadrupled fresh and dried mass of roots and shoot (Table 1 and 2).

There was no significant difference among the AMF treatments for all evaluated parameters regarding the plant vegetative growth. That is, the four AMF species achieved a perfect symbiosis with the host, with nothing affecting the colonization and establishment of their structures in the root system.

The results show a great response of the citrange 'Fepagro C37 Reck' to the presence of AMF in its roots. This is attributed to its root system not having absorbent hairs, a characteristic inherited from one of its parents, Sweet Orange [*Citrus sinensis* (L.) Osbeck]. This orange has few or no root hairs, organs responsible for the absorption of water and nutrients (Zambrosi *et al.*, 2012, Ortas 2012, Wu *et al.*, 2011b).

These beneficial effects of AMF on several plant species are already known and reported in a number of

studies (Costa *et al.*, 2010; Liu & Wu, 2014; Nunes *et al.*, 2011; Wu *et al.*, 2011b), which found that the fresh mass of roots and shoot was greater in mycorrhizal plants than in non-inoculated plants.

Nunes *et al.* (2011) observed in their study that the Okinawa peach rootstock inoculated with AMF *Glomus etunicatum* (Becker & Gerd) kept in a protected environment showed larger diameter, height, leaf area, fresh and dry mass of the root and shoot than non-inoculated plants. In addition, Back *et al.* (2016) evaluated the response of the citrange 'Fepagro C37 Reck' to inoculation with *Scutellospora heterogama* (TH Nicolson & Gerd.), *Gigaspora margarita* (Becker & Hall), *Glomus etunicatum* (Becker & Gerd) and *Acaulospora* sp. and found that the four AMF species provided significant increases in plant height, stem diameter, number of leaves, leaf area, and fresh and dry mass of root and shoot in relation to non-inoculated plants.

Regarding plant nutritional status, Tables 3 and 4 show that AMF also influenced macro and micronutrient levels of the shoot. However, the responses were differentiated, that is, some nutrients had a positive effect, some had negative effect, and others had no effect with inoculation. Among the macronutrients (Table 3), N stands out first, and the relative values were lower than the controls in the plants inoculated with *G. margarita* and *G. etunicatum*. The N contents of the plants inoculated with *S. heterogama* and with *Acaulospora* sp did not differ from the controls.

Table 1: Number of leaves and leaf area (cm²/plant) of the citrus rootstock 'Fepagro C37 Reck' submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	Number of Leaves	Leaf Area (cm ² /plant)
Control	14.08 b	60.536 b
<i>Scutellospora heterogama</i>	27.76 a	209.177 a
<i>Gigaspora margarita</i>	25.16 a	201.880 a
<i>Glomus etunicatum</i>	25.47 a	180.977 a
<i>Acaulospora</i> sp.	24.02 a	175.076 a
CV (%)	14.43	22.01

Means followed by the same letter, between rows, do not differ from each other, by the Duncan test, at the 1% level of significance.

Table 2: Fresh and dry mass of root and shoot (g/plant) of the citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	Shoot (g)		Root (g)	
	FM	DM	FM	DM
Control	2.570 b*	0.770 b**	2.388 b**	0.841 b*
<i>Scutellospora heterogama</i>	9.026 a	3.377 a	10.552 a	3.650 a
<i>Gigaspora margarita</i>	8.314 a	4.180 a	10.920 a	3.867 a
<i>Glomus etunicatum</i>	8.554 a	3.337 a	8.701 a	3.446 a
<i>Acaulospora</i> sp.	7.511 a	2.883 a	8.100 a	3.013 a
CV (%)	21.89	13.24	15.56	16.56

Averages followed by the same letter, between rows, do not differ by Duncan's test, at the level of 5% (*) or 1% (**) of significance.

All AMF-inoculated plants had higher P contents in their tissues compared with controls (Table 3). Among the AMF species, the most efficient in macronutrient accumulation was *Acaulospora* sp., followed by *G. margarita*, *S. heterogama*, and *G. etunicatum*. These results are supported by the literature, since phosphorus is the main nutrient supplied to plants by AMF and it is known for not being mobile in soil and being hardly available to plants (Kiriachek *et al.*, 2009).

The relative values of K in the tissues were only lower in the plants inoculated with *Acaulospora* sp., while the other species were not different from the controls. Ca and S contents were similar between mycorrhizal and control plants. However, the Mg contents were lower in mycorrhizal plants than the controls, independently of the AMF species.

The mycorrhizal response varied according to the nutrient and AMF species for the relative micronutrient values (Table 4). The plants inoculated with *Acaulospora* sp and *G. margarita* had higher percentages of Cu than the other treatments, which were not different. The Zn contents were lower in plants inoculated with *G. margarita*, *G. etunicatum* and *Acaulospora* sp, but plants inoculated with *S. heterogama* were not different from the controls.

The concentration of Fe in the tissues was higher in inoculated plants than in the controls, especially in the

presence of *G. margarita* and *G. etunicatum*. AMF inoculation resulted in a significant dilution effect on the contents of Mn in the tissues when compared with the controls. On the other hand, the levels of B were only inferior in the plants inoculated with *G. margarita*, and no difference was found among the other treatments.

P, Cu, and Fe showed the highest relative values in mycorrhizal plants. Similar results were reported by Nunes *et al.* (2009) and Nunes *et al.* (2013), who found that P contents were higher and Mg were lower in mycorrhizal plants than the controls.

Table 5: Reserve contents of the shoot and root of the citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	Shoot %	Root
Control	33.996 ^{ns}	24.197 ^{ns}
<i>Scutellospora heterogama</i>	32.331	18.601
<i>Gigaspora margarita</i>	32.408	13.886
<i>Glomus etunicatum</i>	33.236	15.702
<i>Acaulospora</i> sp.	31.602	18.317
CV (%)	8.36	20.9

^{ns} Not significant by the Duncan test.

Table 3: Relative content (%) of macronutrients (nitrogen, phosphorus, potassium, calcium, magnesium and sulfur) of the shoot of the citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	N	P	K	Ca	Mg	S
	%					
Control	1.40 a**	0.50 d**	1.33 a*	2.70 ab*	0.29 a**	0.37 ns
<i>Scutellospora heterogama</i>	1.30 ab	0.54 bc	1.37 a	3.03 a	0.25 b	0.33
<i>Gigaspora margarita</i>	1.23 b	0.57 b	1.13 ab	2.60 b	0.23 bc	0.3
<i>Glomus etunicatum</i>	1.16 b	0.53 c	1.09 ab	2.43 b	0.21 c	0.31
<i>Acaulospora</i> sp.	1.43 a	0.65 a	1.00 b	2.67 ab	0.21 c	0.3
CV (%)	3.7	2.86	13.58	5.56	7.12	6.18

Averages followed by the same letter, between rows, do not differ by Duncan's test, at the level of 5% (*) or 1% (**) of significance. ^{ns} Not significant.

Table 4: Relative content (mg/Kg) of micronutrients (copper, zinc, iron, manganese and boron) of the shoot of the citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	Cu	Zn	Fe	Mn	B
	mg/Kg				
Control	131.67 c**	74.00 a*	172.33 c**	405.00** a	48.00 a**
<i>Scutellospora heterogama</i>	134.00 c	71.00 a	230.33 b	150.67 b	47.00 ab
<i>Gigaspora margarita</i>	177.67 b	58.33 b	273.67 a	145.33 b	43.33 b
<i>Glomus etunicatum</i>	144.00 c	54.33 b	291.33 a	139.00 b	45.33 ab
<i>Acaulospora</i> sp.	214.67 a	54.67 b	244.67 b	113.67 c	44.67 ab
CV (%)	3.5	4.61	2.8	2.82	3.2

Averages followed by the same letter, between rows, do not differ by Duncan's test, at the level of 5% (*) or 1% (**) of significance.

Table 6: Percentage of colonization, hyphae density and quantification of arbuscules and vesicles in roots of citrange 'Fepagro C37 Reck' rootstock submitted to the inoculation of four species of arbuscular mycorrhizal fungi (*Scutellospora heterogama*, *Gigaspora margarita*, *Glomus etunicatum* e *Acaulospora* sp.)

Treatment	Colonization (%)	Hyphae	Arbuscules	Vesicles
Control	9.15 b **	0.312 c**	0.271 c**	0.005 b**
<i>Scutellospora heterogama</i>	100.00 a	1.625 b	1.271 b	0.042 b
<i>Gigaspora margarita</i>	100.00 a	2.021 ab	1.958 a	0.500 a
<i>Glomus etunicatum</i>	100.00 a	2.229 a	1.958 a	0.583 a
<i>Acaulospora</i> sp.	100.00 a	2.166 a	1.458 b	0.167 b
CV (%)	0.27	15.66	8.92	21.45

Means followed by the same letter, between rows, do not differ from each other, by the Duncan test, at the 1% level of significance.

Reduction in the nutrient contents caused by AMF are attributed to the effects of dilution of the same in the cells caused by a greater absorption of water. In addition, AMF shows a buffer behavior (Anzanello *et al.*, 2011), that is, in condition of high nutrient availability, they absorb only what is needed, for this reason the results of some nutrients such as Ca, Mg, and Mn had lower values than non-inoculated plants, however, they are already considered high values for plant requirements (SBCS, 2004). Thus, AMF also has the function of regulating the amount of nutrients absorbed, avoiding greater energy expenditure.

AMF had no influence on the reserve contents of the tissues, ranging from 31.6% to 34% in the shoot and 13.9% to 24.2% in the roots (Table 5). The effect of AMF on the reserve content of the plants is variable, and there are reports of positive effects (Nunes *et al.*, 2008a; Nunes *et al.*, 2013), null effect (Nunes *et al.*, 2008b), and even negative effects (Souza *et al.*, 2005) attributed to carbohydrate dilution in the cells as the result of the greater vigor and development provided by the AMF.

As for percentage of colonization, all AMF species colonized the roots of the citrange 'Fepagro C37 Reck' rootstock (Table 6). A small contamination was observed in the control plants. This was probably due to the presence of insect such as ants in the experiment, which may have carried spores to these plants (Table 6). However, contamination has also been reported in other studies (Nunes *et al.*, 2008a; Silveira *et al.*, 2006) and had no interfere in the results because it is considered very low and acceptable.

There was variation among the AMF species regarding the hyphal density, especially *Glomus etunicatum* and *Acaulospora* sp., with high hyphal density, and *Gigaspora margarita* with moderate density (Table 6).

Arbuscules are the AMF structures that exchanges nutrients and photoassimilates with plants (Smith & Smith, 2011). Table 6 shows a considerable amount of arbuscules in the tissues of the roots, characterizing a greater symbiotic activity between the fungus and the plant, explaining the greater vegetative growth of this rootstock.

However, there was a low presence of vesicles in the roots independently of the AMF species (Table 6). Vesicles are globular structures containing granules of glycogen and lipids, thus reserve structures. These structures are formed in periods of low temperature, where there is low activity of the fungus and the host plant (Smith & Smith, 2011). Because the collection of the roots took place in March, at the end of the summer, there was no need for AMF to accumulate reserves. Typically, in southern Brazil, these structures are formed from April/May (autumn months, with reduced temperature and photoperiod) to provide reserves for the AMF during the winter. These findings were corroborated by studies of Nunes *et al.* (2008b) and Nunes *et al.* (2009), which have similar characteristics to this experiment.

CONCLUSIONS

Citrange 'Fepagro C37 Reck' rootstock shows a high response to the presence of AMF in its root system for vegetative grow, independently of the species tested;

AMF influence the relative values of macro and micronutrients in the tissues of citrange 'Fepagro C37 Reck' rootstock, including increased percentage of P, Cu and Fe;

AMF has no effect on the reserve content of citrange 'Fepagro C37 Reck'.

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